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Amendment

In the Specification

Please insert the heading, "Background of the Invention" on page 1, after the title and before line 3.

Please insert the heading, "Brief Summary of the Invention" on page 2, line 3.

Please insert the heading, "Detailed Description of the Invention" on page 2, line 8.

Please insert the heading "Brief Description of the Several Views of the Drawings" on page 20, line 26.

Please replace the paragraph on page 20, lines 24-25, with the following paragraph.

Figure 1 Table A shows individual data for noradrenaline plasma levels which is summarised in Figure 2 1.

Please replace the paragraph on page 20, line 27 to page 21, line 2, with the following paragraph.

Figure 2 1 shows that chronic wasting disorders show increased activity of SNS (sympathetic nervous system) as evidenced by increased plasma noradrenaline levels. All of the cachectic disorders marked (*) have mean plasma noradrenaline levels which are higher than normal. Mean values are given for noradrenaline plasma levels in nmol/1. COPD is chronic occluded pulmonary disease, ncCHF is non-cachectic CHF.

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Please replace the paragraph on page 21, lines 4-7, with the following paragraph.

Figure 3.2 shows that, on average, patients with active wasting disease have 2.5 to 13fold increased aldosterone levels compared to healthy controls (their mean: 43.2 ng/ml, upper limit or normal: 81 ng/ml). Patients with weight loss due to malnutrition have normal aldosterone levels.

Please replace the paragraph on page 21, lines 9-10, with the following paragraph.

Figure [[4]] 3 shows that patients with wasting disease have increased angiotensin II plasma levels. The angiotensin Il plasma levels in various patient types is shown.

Please replace the paragraph on page 21, lines 12-14, with the following paragraph.

Figure 5 4 shows that the frequency of developing cardiac cachexia over time is lower in patients treated with enalapril compared to patients treated with placebo.

Please replace the paragraph on page 27, line 25 to page 28, line 9, with the following paragraph.

We have studied a variety of other cachectic conditions - for instance due to AIDS, liver cirrhosis, chronic obstructive pulmonary disease, chronic renal failure, chronic infections (like pneumonia) and cancer - and we have found activation of the SNS as evidenced by elevated plasma noradrenaline kevels (mean plasma levels were clearly above the upper limit of the normal range, see Figures 1 and 2 Table A and Figure 1). This is not dependent on any specific actiology for the cachectic disorder, in fact we find elevated noradrenaline plasma levels (ic SNS activity) also in cases of idiopathic cachexia, ie cachexia of unknown origin. Nevertheless, we ICH102DraftResponsetoOfficeAction(11_04) ICI 102

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find the activation of the SNS to be specific for eachectic disorders, as it is not seen in patients with a similar degree of weight loss consequent upon malnutrition.

Please insert the following table on page 28, line 23.

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TABLE A

ANOVA Table for HA nmol/I

	DF	Sum of Squares	Mean Square	F-Value	P-Value
Cachexia diag HA-Figure	11	260.240	23.658	2.850	.0020
Residual	103	825.866	8.019		

Model II estimate of between component variance: 1.796 94 cases were omitted due to missing values.

Means Table for HA nmol/I

Effect: Cachexia diag. HA-Figure

AIDS
cachectic CHF
Cancer
chronic renal failure
COPD
healthy controls
ideopathic cachexia
infection
Livercirrh + Cachexia
Malnutrition
more Controls
ne CHF

Count	Mean	Std. Dev.	Std. Err.
6	5.217	4.801	1.960
15	4.870	2.518	.650
2	8.365	5.056	3.575
2	3.686	4.688	3.315
14	3.643	2.305	.616
16	1.940	.687	.172
2	3.835	3.203	2.265
6	6.437	6.966	2.844
6	6.098	5.693	2.324
5	2.967	1.764	.728
3	2.373	1.088	.634
37	2.684	1.344	.221

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Fisher's PLSO for HA nmo// Effect: Cachexia diag.-HA-Figure

Significance Level: 5%

•	Means Diff,	Crit. Diff.	P-Value
AIDS, cachectin CHF	.347	2.718	.8004
AIDS, Cancer	-3.148	4.586	.1768
AIDS, chronic renal failure	1.522	4.586	.5118
AIDS, COPD	1.574	2.740	.2579
AIDS, healthy controls	3.277	2.688	.0174
AIDS, ideopathic cachexia	1.382	4.586	.5514
AIDS, infection	-1.220	3.249	.4572
AIDS, Livercirrh + Cachexia	882	3.243	.5909
AIDS, Malnutrition	2.230	3.249	.1756
AIDS, more Controls	2.643	3.971	.1586
AIDS, nc CHF	2.693	2.472	.0371
cachectic CHF, Cancer	-3.495	4.228	.1042
cachectic CHF, chronic renal failure	1.175	4.226	.5827
cachectic CHF, COPD	1.227	2.087	.2462
cachectic CHF, healthy controls	2.930	2.018	.0049
cachectic CHF, ideopathic cachexia	1.095	4.228	.6283
cachectic CHF, infection	-1.667	2.713	.2547
cachectic CHF, Livercirrh + Cachexia	-1.228	2.713	.3713
cachectic CHF, Malnutrition	-1.869	2.713	.1716
cachectic CHF, more Controls	2.497	3.552	.1663
cachectic CHF, nc CHF	2.286	1.719	.0096
Cancer, chronic renal failure	4.670	5.616	.1022
Cancer, COPD	4.722	4.246	.0296
Cancer, healthy controls	6.425	4.212	.0031
Cancer, ideopathic cachexia			
Cancer, infection	1.928	4.586	.4062
Cancer, Livercirth + Cachexia	2.267	4.586	.3292
Cancer, Malnutrition	5.378	4.586	.0220
Cancer, more Controls	5.992	5.127	.0224
Cancer, nc CHF	5.781	4.077	.0058
chronic renal failure, COPI)	.052	4,246	.9805
chronic renal failure, healthy controls	1.755	4.212	.4105
chronic renal failure, ideopathic cachexia	140	5.516	.9607

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chronic renal failure, infection
chronic renal failure, Livercirth
Cachexia
chronic renal failure, Malnutrition
chronic renal failure, more Controls
chronic renal failure, nc CHF
COPD, healthy controls
COPD, ideopathic cachexia
COPD, Infection
COPD, Livercirrh + Cachexia
COPD, Malnutrition
COPD, more Controls
COPD, ne CHF
healthy controls, ideopathic cachexia
healthy controls, infection
healthy controls, Livercirrh + Cachexis
healthy controls, Malnutrition
healthy controls, more Controls
healthy controls, nc CHF
ideopathic cachexia, infection
ideopathic cachexia, Livercirrh
Cachexia
ideopathic cachexia, Malnutrition
ideopathic cachexia, more Controls
ideopathic cachexia, ne CHF
infection, Livercirrh - Cachexia
infection, Malnutrition
infection, more Controls
infection, ne CHF
Livercirth + Cachexia, Malnutrition Livercirth + Cachexia, more Controls
Livercirrh + Cachexia, no CHF
Malnutrition, more Controls
Malnutrition, no CHF
more Controls, no CHF
more Controls, ne Clar

	-2.742	4.586	.2384
- [-2.403	4.586	.3010
t	.708	4.586	.7600
L	1.322	5.127	.6109
L	1,111	4.077	.5900
L	1.703	2.066	.1085
L	192	4.246	.9285
	-2.794	2.740	.0456
E	-2.456	2.740	.0785
	.856	2.740	.6360
Е	1.269	9.573	.4827
Γ	1.059	1.762	.2362
	-1.895	4.212	.3743
Г	-4.497	2.689	.0013
	-4.158	2.689	.0028
<u> </u>	-1.047	2.689	.4418
Г	433	3.533	.8083
Г	644	1.680	.4491
Γ.	-2.602	4.586	.2631
	-2.263	4.586	.3299
-	.846	4.586	.7144
Γ	1.462	6.127	.5730
Г	1.251	4.077	.5441
	.388	3.243	.8366
ľ	3.450	3.243	.0373
	4.068	3.971	.0450
Г	3.853	2.472	.0026
	3.112	3.243	.0598
Γ	3.725	3.971	.0657
Γ	3.515	2.472	.0058
Г	.613	3.971	.7600
Г	.403	2.472	.7472
Г	210	3.371	.9017

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Please replace the paragraph on page 29, lines 4-14, with the following paragraph.

Aldosterone serum levels have been analysed in a number of subjects with these disorders compared to healthy controls, patients with weight loss due to malnutrition (ie no active wasting disease), and CHF patients without cachexia (see Table below and Figure 3.2). Patients with active wasting disease have on average 2.5 to 13-fold increased aldosterone levels compared to healthy control subjects (their mean: 43.2 ng/ml, upper limit or normal: 81 ng/ml). Patients with weight loss due to malnutrition have normal aldosterone levels. This supports our view that high aldosterone levels are pathophysiologically linked to the presence of chronic active body wasting due, ie cachexia, and that treatment with aldosterone antagonists may be beneficial.

Please delete the paragraphs from page 40, line 21 to page 51, line 3.

Please replace the paragraph on page 51, lines 5-6, with the following paragraph.

Example 10 6: The presence of sympathetic nervous system activation and abnormal sympatho-vagal balance in AIDS - related wasting disease.

Please replace the paragraph on page 53, lines 18-19, with the following paragraph.

Example 11 7: Treatment of a cachectic patient with chronic heart failure with an example beta-blocker (carvedilol).

Please replace the paragraph on page 55, lines 17-18, with the following paragraph.

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Example 12 8: Treatment of cachexia patients with an aldosterone antagonist (spironolactone).